

What is claimed:

1. Isolated nucleic acid having at least 80% nucleic acid sequence identity to a nucleotide
5 sequence comprising the nucleotide sequence shown in Figure 1 (SEQ ID NO:1).
2. Isolated nucleic acid having at least 80% nucleic acid sequence identity to a nucleotide
sequence encoding the polypeptide shown in Figure 2 (SEQ ID NO:2).
- 10 3. Isolated nucleic acid consisting of the full-length coding sequence of the nucleotide
sequence shown in Figure 1 (SEQ ID NO:1).
4. A vector comprising the nucleic acid of Claim 1.
- 15 5. The vector of Claim 4 operably linked to control sequences recognized by a host cell
transformed with the vector.
6. A host cell comprising the vector of Claim 4.
- 20 7. The host cell of Claim 6, wherein said cell is a CHO cell, an *E.coli* cell or a yeast cell.
8. A process for producing a PRO52254 polypeptide comprising culturing the host cell of
Claim 6 under conditions suitable for expression of said PRO52254 polypeptide and recovering said
PRO52254 polypeptide from the cell culture.
- 25 9. An isolated polypeptide having at least 80% amino acid sequence identity to an amino
acid sequence of the polypeptide shown in Figure 2 (SEQ ID NO:2).
10. A chimeric molecule comprising a polypeptide according to Claim 9 fused to a
30 heterologous amino acid sequence.
11. The chimeric molecule of Claim 10, wherein said heterologous amino acid sequence is
an epitope tag sequence or an Fc region of an immunoglobulin.
- 35 12. An antibody which specifically binds to a polypeptide according to Claim 9.
13. The antibody of Claim 12, wherein said antibody is a monoclonal antibody, a humanized
antibody or a single-chain antibody.

14. A composition of matter comprising (a) a polypeptide of Claim 9, (b) an agonist of said polypeptide, (c) an antagonist of said polypeptide, or (d) an antibody that binds to said polypeptide, in combination with a carrier.

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15. The composition of matter of Claim 14, wherein said carrier is a pharmaceutically acceptable carrier.

16. The composition of matter of Claim 14 comprising a therapeutically effective amount of (a), (b), (c) or (d).

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17. An article of manufacture, comprising:

a container;

a label on said container; and

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a composition of matter comprising (a) a polypeptide of Claim 9, (b) an agonist of said polypeptide, (c) an antagonist of said polypeptide, or (d) an antibody that binds to said polypeptide, contained within said container, wherein label on said container indicates that said composition of matter can be used for treating an immune related disease.

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18. A method of treating an immune related disorder in a mammal in need thereof comprising administering to said mammal a therapeutically effective amount of (a) a polypeptide of Claim 9, (b) an agonist of said polypeptide, (c) an antagonist of said polypeptide, or (d) an antibody that binds to said polypeptide.

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19. The method of Claim 18, wherein the immune related disorder is systemic lupus erythematosus, rheumatoid arthritis, osteoarthritis, juvenile chronic arthritis, a spondyloarthropathy, systemic sclerosis, an idiopathic inflammatory myopathy, Sjögren's syndrome, systemic vasculitis, sarcoidosis, autoimmune hemolytic anemia, autoimmune thrombocytopenia, thyroiditis, diabetes mellitus, immune-mediated renal disease, a demyelinating disease of the central or peripheral nervous system, idiopathic demyelinating polyneuropathy, Guillain-Barré syndrome, a chronic inflammatory demyelinating polyneuropathy, a hepatobiliary disease, infectious or autoimmune chronic active hepatitis, primary biliary cirrhosis, granulomatous hepatitis, sclerosing cholangitis, inflammatory bowel disease, gluten-sensitive enteropathy, Whipple's disease, an autoimmune or immune-mediated skin disease, a bullous skin disease, erythema multiforme, contact dermatitis, psoriasis, an allergic disease, asthma, allergic rhinitis, atopic dermatitis, food hypersensitivity, urticaria, an immunologic disease of the lung, eosinophilic pneumonias, idiopathic pulmonary fibrosis, hypersensitivity pneumonitis, a transplantation associated disease, graft rejection or graft-versus-host-disease.

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20. A method for determining the presence of a PRO52254 polypeptide in a sample suspected of containing said polypeptide, said method comprising exposing said sample to an anti-PRO52254 antibody and determining binding of said antibody to a component of said sample.

5 21. A method of diagnosing an immune related disease in a mammal, said method comprising detecting the level of expression of a gene encoding PRO52254 polypeptide (a) in a test sample of tissue cells obtained from the mammal, and (b) in a control sample of known normal tissue cells of the same cell type, wherein a higher or lower level of expression of said gene in the test sample as compared to the control sample is indicative of the presence of an immune related disease in the mammal from which
10 the test tissue cells were obtained.

22. A method of diagnosing an immune related disease in a mammal, said method comprising (a) contacting an anti-PRO52254 antibody with a test sample of tissue cells obtained from said mammal and (b) detecting the formation of a complex between the antibody and the polypeptide in the test
15 sample, wherein formation of said complex is indicative of the presence of an immune related disease in the mammal from which the test tissue cells were obtained.

23. A method of identifying a compound that inhibits the activity of a PRO52254 polypeptide, said method comprising contacting cells which normally respond to said polypeptide with (a)
20 said polypeptide and (b) a candidate compound, and determining the lack responsiveness by said cell to (a).

24. A method of identifying a compound that inhibits the expression of a gene encoding a PRO52254 polypeptide, said method comprising contacting cells which normally express said polypeptide
25 with a candidate compound, and determining the lack of expression said gene.

25. The method of Claim 24, wherein said candidate compound is an antisense nucleic acid.

26. A method of identifying a compound that mimics the activity of a PRO52254
30 polypeptide, said method comprising contacting cells which normally respond to said polypeptide with a candidate compound, and determining the responsiveness by said cell to said candidate compound.

27. A method of stimulating the immune response in a mammal, said method comprising administering to said mammal an effective amount of a PRO52254 polypeptide antagonist, wherein said
35 immune response is stimulated.

28. A method of diagnosing an inflammatory immune response in a mammal, said method comprising detecting the level of expression of a gene encoding PRO52254 polypeptide (a) in a test sample

of tissue cells obtained from the mammal, and (b) in a control sample of known normal tissue cells of the same cell type, wherein a higher or lower level of expression of said gene in the test sample as compared to the control sample is indicative of the presence of an inflammatory immune response in the mammal from which the test tissue cells were obtained.

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29. A method of diagnosing psoriasis in a mammal, said method comprising detecting the level of expression of a gene encoding PRO52254 polypeptide (a) in a test sample of tissue cells obtained from the mammal, and (b) in a control sample of known normal tissue cells of the same cell type, wherein a higher or lower level of expression of said gene in the test sample as compared to the control sample is indicative of the presence of psoriasis in the mammal from which the test tissue cells were obtained.

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30. A method of diagnosing inflammatory bowel disease in a mammal, said method comprising detecting the level of expression of a gene encoding PRO52254 polypeptide (a) in a test sample of tissue cells obtained from the mammal, and (b) in a control sample of known normal tissue cells of the same cell type, wherein a higher or lower level of expression of said gene in the test sample as compared to the control sample is indicative of the presence of inflammatory bowel disease in the mammal from which the test tissue cells were obtained.

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